

**AMENDMENTS TO THE CLAIMS**

**1. (Currently amended)** A method for synthesizing cDNA possessing a ~~consecutive sequence~~ starting with a nucleotide adjacent to a cap structure of mRNA 5'-end nucleotide of (dT)<sub>n</sub>dG, wherein n=0-5, which method comprises the steps of:

(i) annealing a double-stranded DNA primer and an ~~RNA~~mRNA ~~mixture containing mRNA~~ possessing a cap structure,

(ii) preparing a ~~conjugate of an mRNA/cDNA heteroduplex and a double-stranded DNA primer~~ by synthesizing the first-strand cDNA primed with the double-stranded DNA primer using reverse transcriptase, wherein the 3'-end nucleotide of the first-strand cDNA is dC(dA)<sub>n</sub>, wherein n=0-5,

(iii) circularizing ~~the conjugate of the mRNA/cDNA heteroduplex and the double-stranded DNA primer~~ by joining the 3' and 5' ends of the DNA strand containing the first strand cDNA using ligase, and

(iv) replacing the RNA in the mRNA/cDNA heteroduplex with the second-strand cDNA thereby synthesizing the cDNA possessing the 5'-end nucleotide of (dT)<sub>n</sub>dG, wherein n=0-5.

**2. (Currently amended)** The method of claim 1, wherein the mRNA ~~possessing a cap structure~~ is contained in a cell extract.

**3. (Currently amended)** The method of claim 1, wherein the mRNA ~~possessing a cap structure~~ is synthesized by in vitro transcription.

**4. (Currently amended)** The method of claim 1, wherein the primer sequence of the double-stranded DNA primer contains a sequence complementary to a partial sequence of the mRNA ~~possessing a cap structure~~.

**5. (Currently amended)** The method of claim 1, wherein the primer sequence of the double-stranded DNA primer contains an oligo dT complementary to a poly(A) sequence of the mRNA.

~~possessing a cap structure.~~

**6. (Original)** The method of claim 1, wherein the ligase is T4 RNA ligase.

**7. (Currently amended)** The method of claim 1, which comprises the following step between the step (ii) and the step (iii):

(ii') generating a 5'-protruding end or a blunt end at the terminal of the double-stranded DNA primer by cutting ~~the conjugate of the mRNA/cDNA heteroduplex and the double-stranded DNA primer~~ using a restriction enzyme.

**8. (Canceled)**

**9. (Currently amended)** The method of claim 8~~1~~, wherein the double-stranded DNA primer contains a replication origin or both a replication origin and a promoter for cDNA expression.

**10. (Currently amended)** The method of claim 8~~1~~, which further comprises the following step:

(v) incorporating the double-stranded cDNA composed of the first-strand cDNA and the second-strand cDNA into a vector DNA.

**11. (Withdrawn-currently amended)** A cDNA library that is a population of clones containing double-stranded cDNA synthesized by the method of claim 8~~1~~, of which more than 60% of the cDNA clones possesses a 5'-end nucleotide of (dT)<sub>n</sub>dG (n=0-5) followed by a consecutive sequence starting with a nucleotide adjacent to a cap structure of mRNA.

**12. (Canceled)**

**13. (Withdrawn)** A double-stranded DNA primer possessing an oligo (dT)<sub>n</sub> (n=15-100) as a primer part, in which one terminal part of a primer side has an 8-base recognition restriction

enzyme site RE1, and another terminal part has an 8-base recognition restriction enzyme site RE2 and a restriction enzyme site RE3 generating a 5'- protruding end or a blunt end.

**14. (Withdrawn)** The double-stranded DNA primer of claim 13, which contains a replication origin or both a replication origin and a promoter for cDNA expression.

**15. (Withdrawn)** The double-stranded DNA primer of claim 14, which is a vector primer derived from pGCAP10 comprising the nucleotide sequence of SEQ ID NO: 2.

**16. (Withdrawn)** A reagent kit for cDNA synthesis, which comprises the double-stranded DNA primer of claim 14, reverse transcriptase and its reaction buffer solution, T4 RNA ligase and its reaction buffer solution, and model mRNA possessing a cap structure.

**17. (Withdrawn)** A cDNA library that is a population of clones containing double-stranded cDNA synthesized by the method of claim 10, of which more than 60% of the cDNA clones possesses a 5'-end nucleotide of (dT)<sub>n</sub>G (n=0-5) followed by a consecutive sequence starting with a nucleotide adjacent to a cap structure of mRNA.

**18. (Canceled)**

**19. (Withdrawn)** A reagent kit for cDNA synthesis, which comprises the double-stranded DNA primer of claim 15, reverse transcriptase and its reaction buffer solution, T4 RNA ligase and its reaction buffer solution, and model mRNA possessing a cap structure.